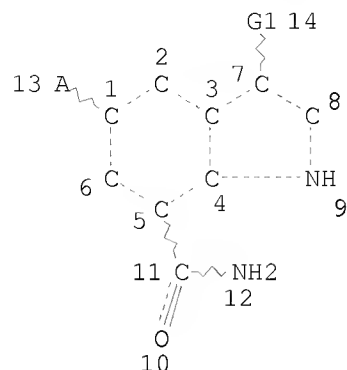


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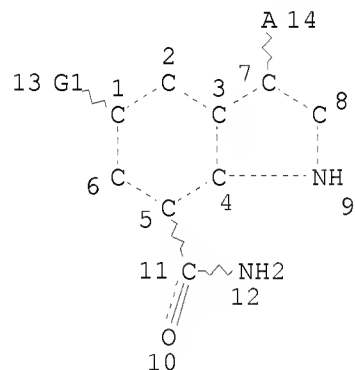
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L5 STR
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DEFAULT ECLEVEL IS LIMITED
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NUMBER OF NODES IS 14
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STEREO ATTRIBUTES: NONE
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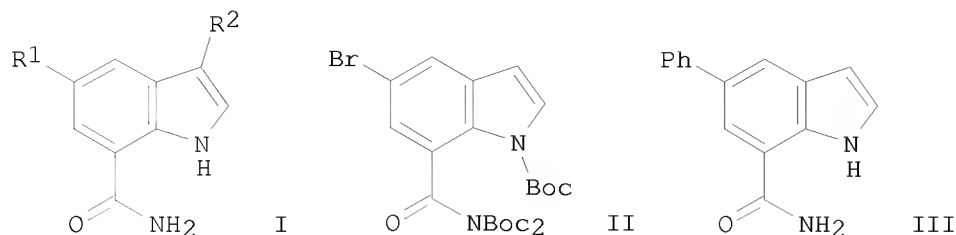
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L14 1 WO2005067923/PN
(WO2005067923/PN)

=> d bib abs

L14 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2009 ACS on STN
AN 2005:673109 CAPLUS
DN 143:172754
TI Preparation of 7-indolecarboxamides as IKK2 kinase inhibitors for the
treatment of such as inflammatory and tissue repair disorders
IN Baldwin, Ian Robert; Bamborough, Paul; Christopher, John Andrew; Kerns,
Jeffrey K.; Longstaff, Timothy; Miller, David Drysdale
PA Smithkline Beecham Corporation, USA
SO PCT Int. Appl., 169 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005067923	A1	20050728	WO 2005-GB85	20050113 <--
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	BR 2005006802	A	20070529	BR 2005-6802	20050113
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	ES 2317184	T3	20090416	ES 2005-701855	20050113
	IN 2006DN03579	A	20070831	IN 2006-DN3579	20060621
	US 20080269200	A1	20081030	US 2006-597154	20060713
	MX 2006008080	A	20060920	MX 2006-8080	20060714
	NO 2006003676	A	20061013	NO 2006-3676	20060815
PRAI	GB 2004-895	A	20040115		
	WO 2005-GB85	W	20050113		
OS	CASREACT 143:172754; MARPAT 143:172754				
GI					



AB Title compds. I [wherein R1, R2 = H, halo, alkylene, alkenylene, (hetero)aryl, etc., and salts, solvates, or physiol. functional derivs. thereof] were prepared as IKK2 kinase inhibitors. For instance, Pd-catalyzed coupling of Boc-protected bromide II (preparation given) with phenylboronic acid followed by deprotection with HCl gave 7-indolecarboxamide III. Most invented compds. were found to have activity >4.8 in the IKK2 assay, in which the degree of phosphorylation of GST-I κ B α was measured as a ratio of specific 665 nm energy transfer signal to reference europium 620 nm signal. Therefore, I and their pharmaceutical compns. are useful in the treatment and prevention of disease states mediated by IKK2 mechanisms, including inflammatory and tissue repair disorders.

RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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ENTER DISPLAY CODE (TI) OR ?:rn
L15 ANALYZE L14 1 RN : 390 TERMS

=> fil reg		
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	ENTRY	SESSION
FULL ESTIMATED COST	20.84	397.62
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-0.82	-0.82

FILE 'REGISTRY' ENTERED AT 10:44:02 ON 13 MAY 2009
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STRUCTURE FILE UPDATES: 12 MAY 2009 HIGHEST RN 1145835-49-9
DICTIONARY FILE UPDATES: 12 MAY 2009 HIGHEST RN 1145835-49-9

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experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

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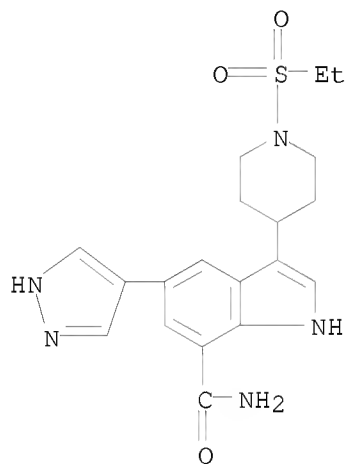
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L16 390 ANSWERS REGISTRY COPYRIGHT 2009 ACS on STN

IN 1H-Indole-7-carboxamide, 3-[1-(ethylsulfonyl)-4-piperidinyl]-5-(1H-pyrazol-4-yl)-

MF C19 H23 N5 O3 S



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

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567770 INDOLE

L17 228 L16 AND INDOLE

=> s l17 and carboximide

4531 CARBOXIMIDE

L18 0 L17 AND CARBOXIMIDE

=> s l17 and carboxamide

2911372 CARBOXAMIDE

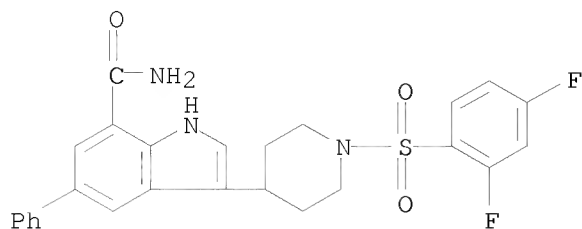
L19 206 L17 AND CARBOXAMIDE

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L19 206 ANSWERS REGISTRY COPYRIGHT 2009 ACS on STN

IN 1H-Indole-7-carboxamide, 3-[1-[(2,4-difluorophenyl)sulfonyl]-4-piperidinyl]-5-phenyl-

MF C26 H23 F2 N3 O3 S

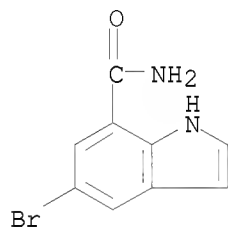


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L19 ANSWER 206 OF 206 REGISTRY COPYRIGHT 2009 ACS on STN
 RN 860624-91-5 REGISTRY
 ED Entered STN: 17 Aug 2005
 CN 1H-Indole-7-carboxamide, 5-bromo- (CA INDEX NAME)
 OTHER NAMES:
 CN 5-Bromo-1H-indole-7-carboxamide
 MF C9 H7 Br N2 O
 SR CA
 LC STN Files: CA, CAPLUS, CASREACT, TOXCENTER, USPATFULL

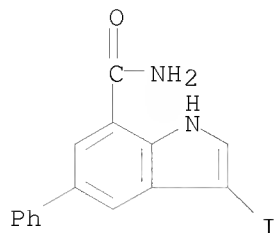


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L19 ANSWER 205 OF 206 REGISTRY COPYRIGHT 2009 ACS on STN
 RN 860624-94-8 REGISTRY
 ED Entered STN: 17 Aug 2005
 CN 1H-Indole-7-carboxamide, 3-iodo-5-phenyl- (CA INDEX NAME)
 MF C15 H11 I N2 O
 SR CA
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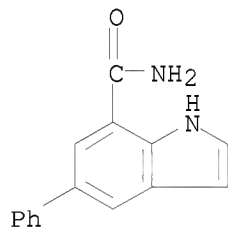


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1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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L19 ANSWER 204 OF 206 REGISTRY COPYRIGHT 2009 ACS on STN
RN 860625-06-5 REGISTRY
ED Entered STN: 17 Aug 2005
CN 1H-Indole-7-carboxamide, 5-phenyl- (CA INDEX NAME)
MF C15 H12 N2 O
SR CA
LC STN Files: CA, CAPLUS, CASREACT, TOXCENTER, USPATFULL



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2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> fil caplus

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FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE	TOTAL
ENTRY	SESSION
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FILE COVERS 1907 - 13 May 2009 VOL 150 ISS 20
FILE LAST UPDATED: 12 May 2009 (20090512/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Feb 2009
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Feb 2009

CAPLUS now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

CAS Information Use Policies apply and are available at:

<http://www.cas.org/legal/infopolicy.html>

This file contains CAS Registry Numbers for easy and accurate

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THE ESTIMATED SEARCH COST FOR 'L15' FOR FILE 'CAPLUS' IS 1,747.20 U.S. DOLLARS

DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N or END:end

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L20 5 L19

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FILE 'REGISTRY' ENTERED AT 10:35:36 ON 13 MAY 2009

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L10	0 S L9 FUL
L11	0 S L5 FUL
L12	STRUC
L13	0 S L12

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FILE 'REGISTRY' ENTERED AT 10:44:02 ON 13 MAY 2009

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L19	206 S L17 AND CARBOXAMIDE

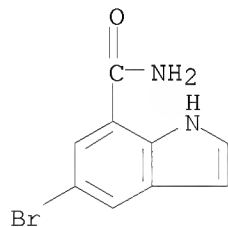
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L21 4 L20 NOT L14

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L21 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN
AN 2008:1180115 CAPLUS
DN 149:425786
TI Preparation of indolecarboxamide derivatives for use as IKK2 inhibitors
IN Boehm, Jeffrey Charles; Busch-Petersen, Jakob; Fu, Wei; Jin, Qi; Kerns, Jeffrey K.; Li, Huijie; Lin, Guoliang; Lin, Xichen; Neipp, Christopher E.
PA Smithkline Beecham Corporation, USA
SO PCT Int. Appl., 245pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2008118724	A1	20081002	WO 2008-US57583	20080320
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PRAI	US 2007-896558P	P	20070323		
OS	MARPAT 149:425786				
IT	860624-91-5P, 5-Bromo-1H-indole-7-carboxamide				
	RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)				
	(preparation of indolecarboxamide derivs. for use as IKK2 inhibitors)				
RN	860624-91-5 CAPLUS				
CN	1H-Indole-7-carboxamide, 5-bromo- (CA INDEX NAME)				



RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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L21 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN

AN 2007:590757 CAPLUS
 DN 147:30940
 TI Preparation of indolecarboxamide derivatives as inhibitors of kinase activity
 IN Kerns, Jeffrey K.; Busch-Petersen, Jakob; Li, Huijie; Boehm, Jeffrey Charles; Nie, Hong; Taggart, John J.
 PA Smithkline Beecham Corporation, USA
 SO PCT Int. Appl., 86pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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OS MARPAT 147:30940

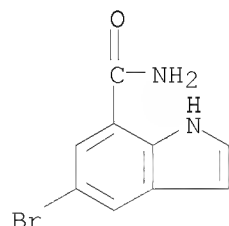
IT 860624-91-5P 860625-19-0P 860625-20-3P
 860625-21-4P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of indolecarboxamide derivs. as inhibitors of kinase activity)

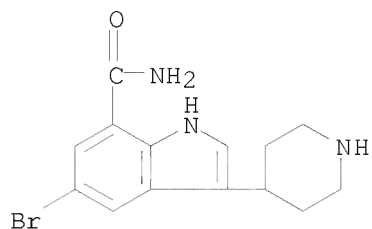
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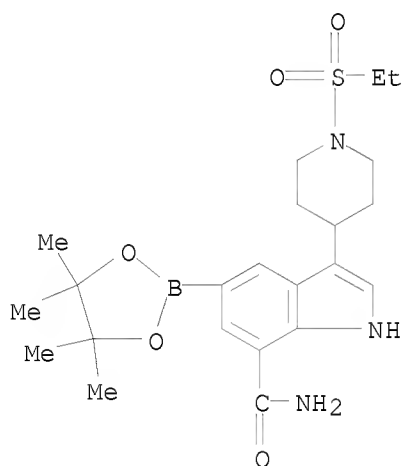


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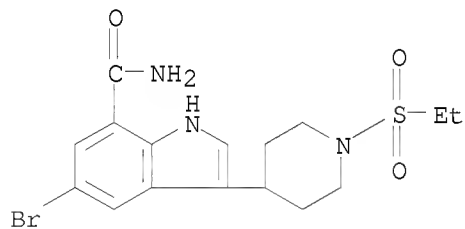
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RN 860625-20-3 CAPLUS
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RN 860625-21-4 CAPLUS
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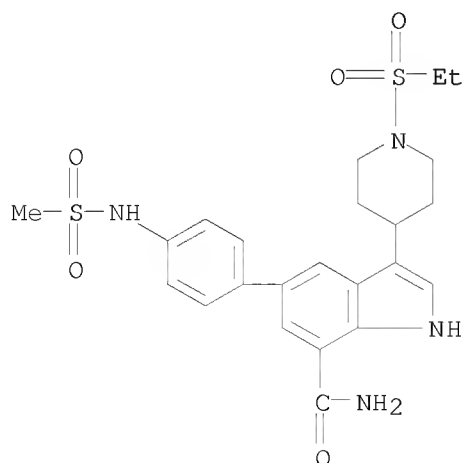
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L21 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2009 ACS on STN
 AN 2007:33976 CAPLUS
 DN 146:142511
 TI Preparation of novel indolecarboxamides as IKK2 inhibitors
 IN Deng, Jianghe; Kerns, Jeffrey K.; Jin, Qi; Lin, Guoliang; Lin, Xichen; Lindenmuth, Michael; Neipp, Christopher E.; Nie, Hong; Thomas, Sonia M.; Widdowson, Katherine L.
 PA Smithkline Beecham Corporation, USA
 SO PCT Int. Appl., 390pp.

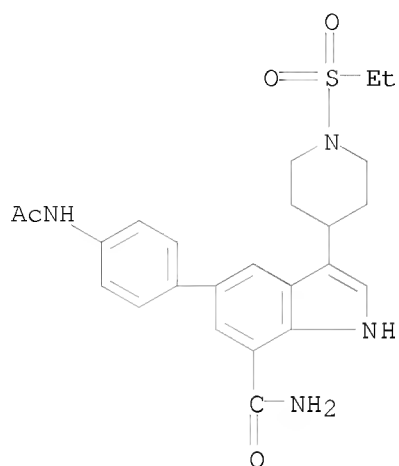
CODEN: PIXXD2

DT Patent
LA English
FAN.CNT 1

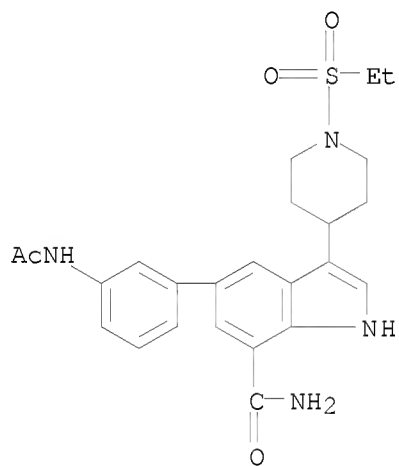
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	IN 2007DN09298	A	20080627	IN 2007-DN9298	20071203
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	KR 2008021077	A	20080306	KR 2007-730656	20071228
	NO 2008000457	A	20080129	NO 2008-457	20080124
	CN 101247804	A	20080820	CN 2006-80030448	20080221
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OS	MARPAT 146:142511				
IT	860626-56-8P 860626-65-9P 860626-67-1P				
	RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)				
	(preparation of novel indolecarboxamides as IKK2 inhibitors)				
RN	860626-56-8 CAPLUS				
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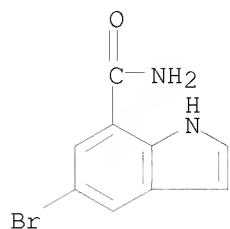
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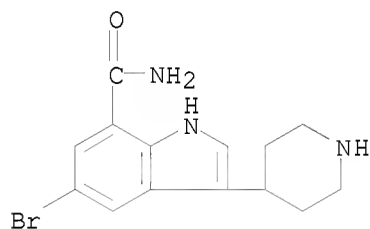
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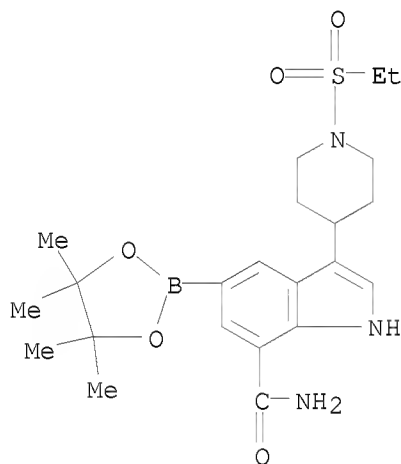
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 RN 860624-91-5 CAPLUS
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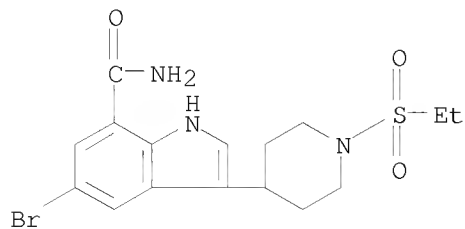
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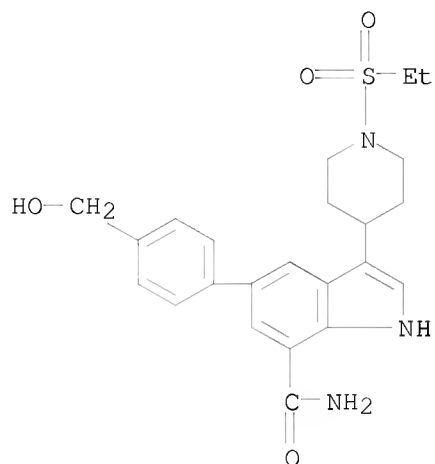
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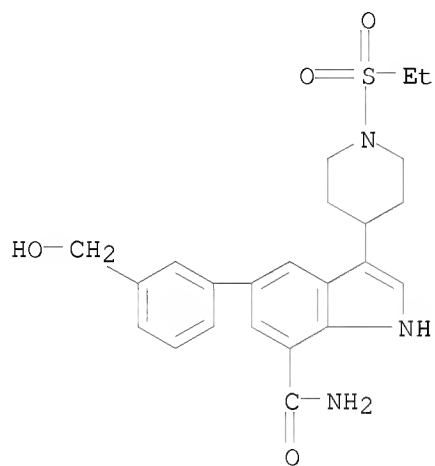
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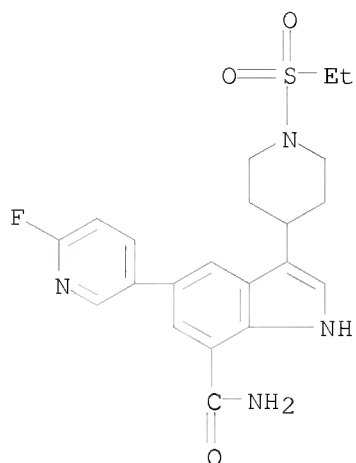
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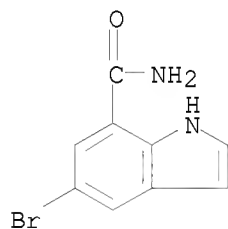
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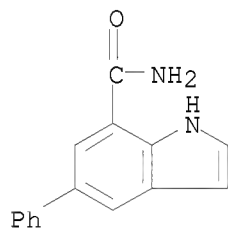
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 DN 144:350542
 TI Indole derivatives as IKK2 inhibitors and their preparations,
 pharmaceutical compositions, and use for treatment of diseases associated
 with inappropriate IKK2 activity such as rheumatoid arthritis, asthma and
 chronic obstructive pulmonary disease
 IN Kerns, Jeffrey K.; Lindenmuth, Michael; Lin, Xichen; Nie, Hong; Thomas,
 Sonia M.
 PA Glaxo Group Limited, UK
 SO PCT Int. Appl., 220 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006034317	A2	20060330	WO 2005-US33752	20050921
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	EP 1793826	A2	20070613	EP 2005-798511	20050921
	R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, YU			
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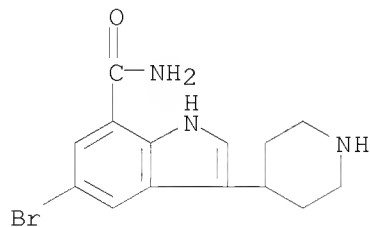
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	US 2005-695454P	P	20050630		
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	860626-15-9P 860626-16-0P				
	RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)				
	(intermediate; preparation of indole derivs. as IKK2 inhibitors and for treatment of diseases associated with inappropriate IKK2 activity such as rheumatoid arthritis, asthma and chronic obstructive pulmonary disease)				
RN	860624-91-5 CAPLUS				
CN	1H-Indole-7-carboxamide, 5-bromo- (CA INDEX NAME)				



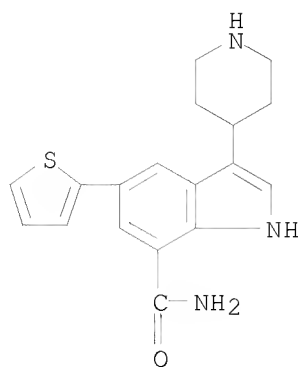
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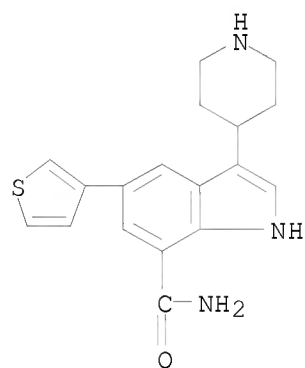
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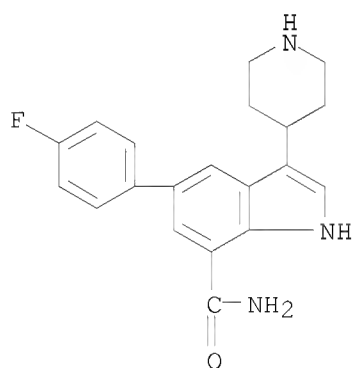
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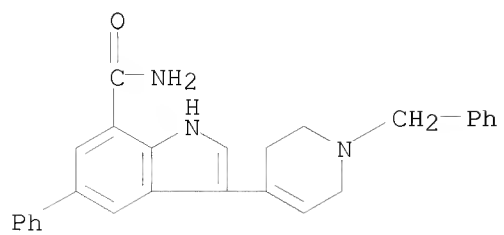
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RN 860625-44-1 CAPLUS
 CN 1H-Indole-7-carboxamide, 5-(4-fluorophenyl)-3-(4-piperidinyl)- (CA INDEX NAME)



RN 860626-15-9 CAPLUS
 CN 1H-Indole-7-carboxamide, 5-phenyl-3-[1,2,3,6-tetrahydro-1-(phenylmethyl)-4-pyridinyl]- (CA INDEX NAME)



RN 860626-16-0 CAPLUS

CN 1H-Indole-7-carboxamide, 5-phenyl-3-(4-phenylpiperidin-1-yl)- (CA INDEX NAME)

